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L20
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L22
               STRUCTURE UPLOADED
L23
            2 S L19
L24
             0 S L20
             0 S L21
L25
L26
             2 S L22
    FILE 'CAPLUS' ENTERED AT 18:37:37 ON 03 APR 2005
L27
            36 S L23
L28
             1 S L26
L29
       306586 S IMAGE OR IMAGING
            5 S L29 AND (L27 OR L28)
L30
L31
            31 S L27 NOT L30
      244973 S PHOTOSENSITIVE OR PHOTOACTIVATABLE OR RECORDING OR PHOTOPOLYM
L32
L33
            9 S L31 AND L32
L34
           114 S 5MW OR 5 MILLIWATTS OR 5MILLI WATTS OR 5 MILLI WATTS
L35
           145 S 2MW OR 2 MILLIWATTS OR 2MILLI WATTS OR 2 MILLI WATTS
            50 S 3MW OR 3 MILLIWATTS OR 3MILLI WATTS OR 3 MILLI WATTS
L36
L37
            43 S 4MW OR 4MILLIWATTS OR 4MILLI WATTS OR 4 MILLI WATTS
L38
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             0 S L34 AND L27
L39
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             0 S L26
L41
L42
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L43
             2 S L42 AND L40
    FILE 'CAPLUS' ENTERED AT 18:48:03 ON 03 APR 2005
    65395 S MW OR MILLIWATTS OR MILLI WATTS
L44
L45 .
         0 S L44 AND (L27 OR L28)
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L30 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1991:502924 CAPLUS

DN 115:102924

TI Recording material containing unsaturated ketone and electron acceptor

IN Satomura, Masato; Takashima, Masanobu; Sano, Masajiro; Yanagihara, Naoto

PA Fuji Photo Film Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

111111111111111111111111111111111111111				
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 03049982	A2	19910304	JP 1989-186249	19890719
DDAT .TD 1989_186249		19990719		

The material uses color formation in contacting an electron acceptor with an unsatd. ketone obtained by condensation with substituted amino-containing aldehyde and ketone. The material is used in pressure-sensitive recording and thermal recording. A material containing unsatd. ketone condensed with melamine formaldehyde and I and Zn 3,5-bismethylbenzylsalicylate gave a clear and hard image.

IT 19226-99-4

RL: USES (Uses)

(coloring agent, pressure-sensitive recording material containing)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L30 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1990:100741 CAPLUS

DN 112:100741

TI Photopolymerization initiator and thermal-transfer recording medium

IN Okuma, Norio

PA Canon K. K., Japan; Sanyo Chemical Industries Ltd.

SO Jpn. Kokai Tokkyo Koho, 18 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 01174502	A2	19890711	JP 1987-335731	19871228
PRAI JP 1987-335731		19871228		

AB The photopolymn. initiator is composed of I or II [Arl, Ar2 = aromatic ring, heterocyclic ring; R1 = H, C1-10 alkyl, alkenyl, alkoxy, or alkylthio, C6-12 aryl, aryloxy, or heterocyclic ring with number of C and non-C atoms to be 5-15; X = non-metallic atom for forming a ring], and III [Y = halogen; R = alkyl, aryl, alkenyl; Q = CY3, NH2, etc.]. The thermal-transfer recording layer is composed of the photoinitiator, and monomer, oligomer, or polymer with unsatd. double bond or these mixture An image -forming material may be encapsulated. This initiator is especially useful in one-shot color recording.

IT 19226-99-4

RL: USES (Uses)

(photopolymn. initiator composition containing triazine derivative and)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L30 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1988:560691 CAPLUS

DN 109:160691

TI Visible laser-sensitive photoimaging compositions and processes

IN Tamaoki, Nobuyuki

PA Toyobo Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	JP 63055539	A2	19880310	JP 1986-126823	19860530
PP A T	TP 1986-126823		19860530		

OS MARPAT 109:160691

AB The title material contains (a) a photooxidn. agent which becomes an oxidation agent by irradiation of a visible laser (420-550 nm), (b) a leuco body

which becomes a dye by reacting with the oxidation agent, (c) a photopolymn. initiator which generates a radical with the dye by irradiating with light (550-700 nm), and (d) >1 ethylenically unsatd. compound nongaseous at room temperature. The method for <code>image</code> formation involves irradiating with a visible laser 420-550 nm, and then with light 550-700 nm to harden <code>image</code> areas. The material shows high sensitivity to visible laser.

IT 19226-99-4

RL: USES (Uses)

(visible laser-sensitive image forming material containing)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L30 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1985:496389 CAPLUS

DN 103:96389

ΤI Photoinsolubilizing resin composition

Agency of Industrial Sciences and Technology, Japan PA

Jpn. Kokai Tokkyo Koho, 4 pp. SO

CODEN: JKXXAF

DT Patent

T.A Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	~~				
ΡI	JP 60078443	A2	19850504	JP 1983-186398	19831005
	JP 05065869	B4	19930920		
PRAI	JP 1983-186398		19831005		

Resin composition contains ethylenic monomer and a photoinitiator, which is a AB combination of an unsatd. ketone having the general formula I (R, R1, R4, R5 = alkyl; R2, R3 = H, or are combined to form C1-3 alkylene group that is a part of a ring system; n = 0,1) with a diaryliodonium salt. The unsatd. ketone effectively promotes photodecompn. of the iodonium compound, and the use of the photoinitiator provides high sensitivity of the resin composition, especially at longer wavelengths. Thus, 0.01 part of diphenyliodonium

hexafluorophosphate and 0.01 part of bis(p-dimethylaminobenzylidene)aceton e were added to 1 part of 10% dioxane solution of a copolymer prepared by introducing methacryloyl group to 1:1 chloromethylstyrene-Me methacrylate copolymer, and the mixture was coated on an anodized Al plate. Photosensitivity to Xe lamp radiation was 32 times higher than that of com. products.

IT 19226-99-4

RL: USES (Uses)

(photoimaging resin composition containing)

RN 19226-99-4 CAPLUS

Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) CNINDEX NAME)

L30 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN

1985:176527 CAPLUS

DN 102:176527

Photographic recording using photohardenable materials TI

IN Grossa, Mario

Du Pont de Nemours (Deutschland) G.m.b.H., Fed. Rep. Ger. PA

Ger., 8 pp. SO CODEN: GWXXAW

DΤ Patent LΑ German

FAN.CNT 1

P

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	DE 3335309	C1	19840816	DE 1983-3335309	19830929
	US 4595651	Α	19860617	US 1984-647809	19840906
PRAI	DE 1983-3335309	A	19830929		

AB A process for the photoimaging of selective regions of an original as equi-d. or contour images involves imagewise exposing a photohardenable material to light of wavelength A which desensitizes the material to exposure by light of wavelength B, and then photohardening the material by exposure to light of wavelength B. The <code>image</code> formation proceeds through ≥ 2 imagewise exposures with light of different wavelengths and addnl. nonimagewise, photohardening final exposure. Thus, a PET support was overcoated with a CH2Cl2 solution containing poly(vinyl acetate) 12.6, poly(Me methacrylate) 31.1, trimethylolpropane triacrylate 35.6, oxyethylated trimethylolpropane triacrylate 8.0, oxyethylated hexadecanol 8.0, 2-(o-chlorophenyl)-3,4-diphenylimidazole 1.6, 2-mercaptobenzoxazole 08, an inhibitor precursor 2.0, and a sensitizer 0.3% and then laminated at 100° to a white paper. The laminate was then exposed through a halftone wedge (wedge constant of $\sqrt{2}$) for 10 s to light of λ = 400-700 nm and 90 s to light of λ = 300-400 nm followed by a nonimage exposure for 15 s to light of λ = 400-700 nm. After the exposure the polyester layer was stripped off, and the <code>image</code> toned. The width of the equi-d. was 1.7.

IT 19226-99-4

RL: USES (Uses)

(photoimaging compns. containing, for contour or equidensity images) 19226-99-4 CAPLUS

RN

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L33 ANSWER 1 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN
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AN 1995:389593 CAPLUS

DN 122:147301

TI Distinguishing markable photoresist material

IN Grossa, Mario

PA Du Pont de Nemours (Deutschland) GmbH, Germany

SO Ger. Offen., 6 pp. CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	DE 4240141	A1	19940601	DE 1992-4240141	19921128
PRAI	DE 1992-4240141		19921128		

The title material comprises: (1) a polymer binder; (2) an ethylenically unsatd. addition polymerizable compound; (3) a leuco dye; (4) a UV light-absorbing free radical-forming hexaaryl bisimidazole-type initiator; (5) a spectral sensitizer dye for visible or IR region; and (6) an another free radical-forming initiator of the type aromatic carbonyl compound where the carbonyl compound absorbs only in the UV region and the spectra sensitizer selectively sensitizes the bisimidazole-type initiator. The composition produces high d. images.

IT 19226-99-4

RL: MOA (Modifier or additive use); USES (Uses) (carbonyl compound as photopolymn. initiator)

RN 19226-99-4 CAPLUS

L33 ANSWER 2 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1995:74750 CAPLUS

DN 122:146976

TI Visible light **photopolymerization** initiated by hexaarylbiimidazole (HABI's)

AU Yang, Weidong; Yang, Yongyuan; Wang, Junshen; Zhang, Cunlin; Yu, Meiwen

CS Institute of Photographic Chemistry, Academia Sinica, Beijing, Peop. Rep. China

SO Journal of Photopolymer Science and Technology (1994), 7(1), 187-92 CODEN: JSTEEW; ISSN: 0914-9244

DT Journal

LA English

AB Kinetics of Me methacrylate (MMA) photopolymn. was studied using 2,5-bis(4'-diethylaminobenzylidene)cyclopentanone (DEAP) and 2,5-bis(4'-dimethylaminobenzylidene)cyclopentenone (DMAP) as visible light sensitizers and 2-chlorohexaarylbiimidazole (o-Cl-HABI) as polymerization initiator. The photopolymn. study were carried out in toluene solution at 30°C. The polymerization rate was proportional to the concentration with exponent of 0.85, -0.28, 0.30, 1.0 for DEAP, o-Cl-HABI, 3-mercapto-4-methyl-4H-1,2,4-triazole additive,MMA, resp. The photopolymer materials can record stable hologram with sensitivity of 30.apprx.300mJ/cm2 and resolution of 4000 1/mm.

IT 19226-99-4

RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)

(photoinitiator; kinetics of visible light-induced photopolymn. of Me methacrylate initiated by hexaarylbiimidazole for holog.

recording)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L33 ANSWER 3 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1993:437581 CAPLUS

DN 119:37581

TI Photopolymerizable compositions for lithographic printing plates

IN Okamoto, Hiroaki

PA Okamoto Kagaku Kogyo Kk, Japan

SO Jpn. Kokai Tokkyo Koho, 10 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN. CNT 1

1741. CIVI 1				
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI JP 05078410	A2	19930330	JP 1991-268302	19910919
PRAI JP 1991-268302		19910919		

OS MARPAT 119:37581

AB The title compns. comprise radical polymerizable compds. having ≥ 2 ethylenically unsatd. double bonds and photoinitiator mixts. containing p-RR1NC6H4(CH:CH)nCH:CR2COCR22:CH(CH:CH)np-C6H4NRR1 [R, R1 = C1-6 alkyl, cycloalkyl, hydroxyalkyl; RR1 may be tetramethylethylene (sic), pentamethylethylene (sic), oxybisethylene; R2, R22 = H, alkyl, Ph; R2R22 may be (CH2)2, (CH2)3, or CO; n = 0-3], PPh3 and/or quaternary phosphonium salts, thiols I (Z = NH, S, O), and tetrazolium derivs. II [R3-5 = alkyl, (un)substituted aryl, styryl, thienyl, trimethylammoniumindolyl; X = Cl, perchloride]. The compns. show high Ar laser sensitivity and are useful as **photosensitive** layer in presensitized lithog. printing plates.

IT 19226-99-4

RL: USES (Uses)

(photoinitiators containing, in **photosensitive** layers for lithog. printing plates)

RN 19226-99-4 CAPLUS

L33 ANSWER 4 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1993:244666 CAPLUS

DN 118:244666

TI Photosensitive resin composition and hologram recording media and its recording

IN Kobayashi, Tatsu; Yoshinaga, Yoko

PA Canon Kk, Japan

SO Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	JP 04368948	A2	19921221	JP 1991-171953	19910618
DDAT	TP 1991-171953		19910618		

OS MARPAT 118:244666

AB The composition contains a vinylcarbazole-containing polymer and a halo-containing Si

compound R1R2R3R4Si [R2-4 = halo, H, (substituted) alkyl, cycloalkyl, OH, alkoxy, (substituted) aryl, aryloxy, alkylcarbonyl, alkoxycarbonyl, NH2, dialkylamino, NO2, CN; R1 and R2 may form a ring with Si; R1 = halo] as a crosslinking agent. The media contain the composition The method exposuring the media by an UV ray or a visible light.

IT 19226-99-4

RL: USES (Uses)

(photosensitized pigment, **photosensitive** resin containing, for hologram **recording**)

RN 19226-99-4 CAPLUS

L33 ANSWER 5 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1991:33174 CAPLUS

DN 114:33174

TI Cycloalkene compounds for thermal recording materials

IN Wakasugi, Kazuyuki; Kikkawa, Katsumasa; Yamaguchi, Masahiko; Motohashi, Katsuichi

PA Hodogaya Chemical Co., Ltd., Japan

SO Eur. Pat. Appl., 14 pp.

CODEN: EPXXDW

DT Patent

LA English

FAN. CNT 1

FAN.	CNT I				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	EP 364120	A2	19900418	EP 1989-309647	19890921
	EP 364120	A3	19910814		
	EP 364120	B1	19930811		
	R: DE, FR, GB				
	JP 02103265	A2	19900416	JP 1988-255920	19881013
	JP 2566635	B2	19961225		

US 4987262 A 19910122 US 1989-412127 19890925

PRAI JP 1988-255920 A 19881013

OS MARPAT 114:33174

AB The title compds. have the general formula I [R1, R2, R6 = C1-4 alkyl; R3 = H, R1; R4 and R5 together form -CH2CH2- or -(CH2)3-] and are used as color formers in thermal **recording** materials.

IT 19226-99-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation and reaction of, in preparing color former for thermal recording material)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L33 ANSWER 6 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1990:554275 CAPLUS

DN 113:154275

TI Cycloalkene compounds and **recording** materials using these compounds as color formers

IN Wakasugi, Kazuyuki; Yamaguchi, Masahiko; Sato, Hiroko; Motohashi, Katsuichi

PA Hodogaya Chemical Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 12 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

1111. C1	11 1				
F	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
_					
PI J	JP 02135264	A2	19900524	JP 1988-288768	19881117
PRAI J	JP 1988-288768		19881117		

OS MARPAT 113:154275

The title color formers I (R1, R2 = lower alkyl; R3 = H, lower alkyl; R4, R5 = (CH2)1-3; R6 = lower alkyl, substituted phenyl], showing strong absorption in the near-IR region when developed, are prepared Thus, 4-Me2NC6H4CHO was treated with cyclohexanone in EtOH in the presence of NaOH to give 2,6-bis[4-(dimethylamino)benzylidene)cyclohexanone which was reduced with NaBH4 in tetrahydrofurfuryl alc.-THF and treated with HClO4 to give 2,6-bis[4-(dimethylamino)benzylidene]cyclohexanium perchlorate which was then treated with PhSO2Na.2H2O in MeOH to give I [R1 = R2 = Me, R3 = H, R4R5 = (CH2)3, R6 = Ph], greenish blue or bluish green on clay, resin, salicylates, etc.

IT 19226-99-4P

RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)

(manufacture and reduction of)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L33 ANSWER 7 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1990:88367 CAPLUS

DN 112:88367

TI Visible ray-sensitive **photopolymerizable** vinyl polymer compositions

IN Imahashi, Satoshi; Saito, Atsushi

PA Toyobo Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE
PI JP 01205153 A2 19890817 JP 1988-29581 19880210

PRAI JP 1988-29581 19880210

The title composition contains ≥1 vinyl compds. nongaseous at room temperature, an organic peroxide group ArC(:0)O2- [Ar = Ph (substituted with ≥1 groups selected from Ph, amino, carbonyl, and halo), C1-4 alkyl, C1-4 alkoxy]-containing compound, and p-aminophenyl unsatd. ketone I [R1-2 = H, C1-5 alkyl; R3 = methylidine, C1-5 alkylenylidine forming ring with CO; R4 = C, (substituted) Ph, a group forming indanone or tetralone with R3 and CO; R5 = p-R6R7NC6H4(CH:CH)nCH:; R6-7 = H, C1-5 alkyl; m, n = 0, 1]. The composition is useful for a **photoresist** or a printing plate. Thus, a transparent PET film was coated with a composition of methacrylic acid-Me methacrylate copolymer, tetraethylene glycol diacrylate, 3,3',4,4'-tetra(tert-butylperoxycarbonyl)benzophenone, 2,5-bis(4'-diethylaminobenzylidene)cyclopentanone, MeOH, and EtOAc, dried, coated with aqueous poly(vinyl alc.), dried, neg. pattern-wise irradiated at

490 nm, aqueous Na2CO3-developed, and washed to give a highly-cured pattern.

IT 19226-99-4

RL: USES (Uses)

(photoresists from, visible ray-sensitive)

RN 19226-99-4 CAPLUS

L33 ANSWER 8 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1989:544118 CAPLUS

DN 111:144118

TI Photopolymerization initiator and photoresist composition containing same

IN Kaji, Makoto; Kaneko, Futami; Hayashi, Nobuyuki

PA Hitachi Chemical Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DT Patent LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	JP 63309502	A2	19881216	JP 1987-144902	19870610
	JP 2536528	B2	19960918		
PRAI	JP 1987-144902		19870610		

AB The title photopolymn. initiator contain (I) [R1-5 = H, C1-12 alkyl, halo; R6 = H, C1-12 alkyl, cycloalkyl, C1-12 hydroxyalkyl, C1-12 alkoxyalkyl, C1-12 aminoalkyl, aryl; R7-8 = H, C1-8 alkyl] and (II) [R9,10 = C1-9 alkyl; R11-14 = H, C1-4 alkyl, C1-4 alkoxy, halo; R15, R16 = H, C1-4 alkyl; Z = aryl, styryl], and the title **photoresist** composition contains I, II, and an addition polymerizable compound with b.p. ≥100°. High sensitivity to visible light is confirmed by the photopolymn. initiator.

IT 19226-99-4

RL: USES (Uses)

(photopolymn. initiator containing, for photoresist composition)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L33 ANSWER 9 OF 9 CAPLUS COPYRIGHT 2005 ACS on STN

AN 1972:427427 CAPLUS

DN 77:27427

TI Photoactivatable compositions

IN Baum, Martin D.; Henry, Cyrus P., Jr.

PA du Pont de Nemours, E. I., and Co.

SO Ger. Offen., 44 pp.

CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

PAIN. CIVI I				
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI DE 2133515	Α	19720113	DE 1971-2133515	19710706
DE 2133515	B2	19760812		
DE 2133515	C3	19770324		
US 3652275	Α	19720328	US 1970-53686	19700709
FR 2100341	A 5	19720317	FR 1971-24823	19710707
GB 1341244	Α	19731219	GB 1971-31879	19710707
BE 769695	A1	19720110	BE 1971-105645	19710708
PRAI US 1970-53686	Α	19700709		

Photoactivatable compns. that can be utilized as light filters or photooxidn. or photopolymn. initiators are composed of a hexaarylbiimidazole whose principal radiation absorption bands are in the uv region of the spectrum and which dissocs. into triarylimidazolyl radicals on uv irradiation, and a sensitizing bis(p-aminophenyl)-α,β-unsatd. ketone (I), where R1 and R2 are alkyl or H; R3 is H, alkyl, C1, or MeO; R4 and R5 are H, alkyl, or R4R5 is CH2CH2,

CH2CH2CH2, or CH2CH2CH2CH2; n=0, 1, and in which I has its main absorption bands in the visible region. Thus, a Mylar film coated with a solution containing cellulose acetate butyrate 13.2, 2,2'-bis(o-chlorophenyl)-4,4',-5,5'-tetrakis(m-methoxyphenyl)biimidazole 3, 2-mercaptobenzoxazole 0.1, I (R1 and R2 are Et, R3 is H, (R4R5) is CH2CH2, n=0) (II) 0.05 g, and triethylene glycol dimethacrylate 12.5 ml is covered with a polyester film and exposed to filtered radiation of 366 and 430 nm. The exposure time for complete photopolymn. for irradiation at 366 and 430 nm is 2 and 2 sec, resp., vs. 8 and 2 sec, resp., for a II-free control.

IT 19226-99-4

RL: USES (Uses)

(photosensitizer, for **photoactivatable** compns. containing hexaarylbiimidazole)

RN 19226-99-4 CAPLUS

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

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FILE 'CAPLUS' ENTERED AT 18:37:37 ON 03 APR 2005
L27
            36 $ L23
L28
             1 S L26
       306586 S IMAGE OR IMAGING
L29
L30
            5 S L29 AND (L27 OR L28)
            31 S L27 NOT L30
L31
        244973 S PHOTOSENSITIVE OR PHOTOACTIVATABLE OR RECORDING OR PHOTOPOLYM
L32
            9 S L31 AND L32
L33
           114 S 5MW OR 5 MILLIWATTS OR 5MILLI WATTS OR 5 MILLI WATTS
L34
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L35
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L36
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L37
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            0 (L34 OR L35 OR L36 OR L37) AND (L27 OR L28 OR L33 OR L31)
=> s 134 and 127
L39
        0 L34 AND L27
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L40 ANSWER 1 OF 4 USPATFULL on STN
       91:7159 USPATFULL
ΑN
ΤI
       Cycloalkene compounds useful in recording materials
IN
       Kazuyuki, Wakasugi, Tokyo, Japan
       Kikkawa, Katsumasa, Tokyo, Japan
       Yamaquchi, Masahiko, Tokyo, Japan
      Motohashi, Katsuichi, Tokyo, Japan
       Hodogaya Chemical Co., Ltd., Tokyo, Japan (non-U.S. corporation)
PA
PΤ
      US 4987262
                               19910122
      US 1989-412127
ΑI
                               19890925 (7)
      JP 1988-255920
PRAI
                           19881013
DT
      Utility
FS
      Granted
EXNAM
      Primary Examiner: Raymond, Richard L.
LREP
       Sherman and Shalloway
CLMN
      Number of Claims: 3
ECL
      Exemplary Claim: 1
DRWN
      No Drawings
LN.CNT 400
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      A novel cycloalkene compound represented by the following general
       formula [1]: ##STR1## wherein each of R.sup.1 and R.sup.2 is a C.sub.1
       -C.sub.4 alkyl group, R.sup.3 is a hydrogen atom , or a C.sub.1 -C.sub.4
      alkyl group, R.sup.4, together with R.sup.5, forms --CH.sub.2 --CH.sub.2
       --or --CH.sub.2 --CH.sub.2 --, and R.sup.6 is a C.sub.1
       -C.sub.4 alkyl group.
IT
   19226-99-4P
        (preparation and reaction of, in preparing color former for thermal
recording
        material)
RN
     19226-99-4 USPATFULL
CN
     Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI)
       INDEX NAME)
```

L40 ANSWER 2 OF 4 USPATFULL on STN

AN 86:35649 USPATFULL

TI Process for producing equidensity images using photohardenable materials

IN Grossa, Mario, Dreieich, Germany, Federal Republic of

PA E. I. Du Pont de Nemours and Company, Wilmington, DE, United States

(U.S. corporation)

PI US 4595651 19860617 AI US 1984-647809 19840906 (6)

PRAI DE 1983-3335309 19830929

DT Utility FS Granted

EXNAM Primary Examiner: Kittle, John E.; Assistant Examiner: Hamilton, Cynthia

CLMN Number of Claims: 6 ECL Exemplary Claim: 1,2

DRWN 2 Drawing Figure(s); 2 Drawing Page(s)

LN.CNT 460

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Process for photographic recording of selective areas of an original

image, e.g., as equidensities and/or contour images, on a photohardenable material, e.g., (a) nongaseous monomer compound, (b) organic photoinitiator or photoinitiator/sensitizer system and (c) photoinhibitor activatable by ultraviolet radiation, which comprises in either order (A) exposing the photohardenable layer with desensitizing radiation, and (B) photohardening the layer with photohardening radiation, exposures (A) and (B) being imagewise exposures with the proviso that if exposure (B) is the first imagewise exposure, a nonimagewise final exposure is utilized.

IT 19226-99-4

RN

(photoimaging compns. containing, for contour or equidensity images) 19226-99-4 USPATFULL

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) (CA INDEX NAME)

L40 ANSWER 3 OF 4 USPATFULL on STN

AN 74:62063 USPATFULL

TI ARYLIDENE CYCLANONES IN INHIBITING ANDROGEN ACTION

IN Scanlon, William B., Indianapolis, IN, United States

PA Eli Lilly and Company, Indianapolis, IN, United States (U.S.

corporation)

PI US 3857953 19741231

AI US 1972-315947 19721218 (5)

RLI Continuation-in-part of Ser. No. US 1971-182123, filed on 20 Sep 1971, now abandoned

DT Utility

FS Granted

EXNAM Primary Examiner: Friedman, Stanley J.

LREP Martens, Jr., William C., Smith, Everet F.

CLMN Number of Claims: 7 ECL Exemplary Claim: 1

DRWN No Drawings

LN.CNT 581

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method for inhibiting action of androgens by administering an effective amount of an arylidene cyclanone, anti-androgen compositions containing an arylidene cyclanone, and selected arylidene cyclanones.

IT 19226-99-4P

(preparation of)

RN 19226-99-4 USPATFULL

L40 ANSWER 4 OF 4 USPATFULL on STN

AN 72:16254 USPATFULL

TI HEXAARYLBIIMIDAZOLE BIS (p-DIALKYL-AMINOPHENYL- α , β -

UNSATURATED) KETONE COMPOSITIONS

IN Baum, Martin D., Wilmington, DE, United States

Henry, Jr., Cyrus P., Wilmington, DE, United States

PA E. I du Pont de Nemours and Company, Wilmington, DE, United States

PI US 3652275 19720328

AI US 1970-53686 19700709 (5)

DT Utility

FS Granted

EXNAM Primary Examiner: Torchin, Norman G.; Assistant Examiner: Fichter,

Richard E.

LREP Powell; John R.

CLMN Number of Claims: 36

DRWN No Drawings

LN.CNT 1249

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Compositions comprising a hexaarylbiimidazole and a selected bis(p-aminophenyl-... α , β -unsaturated) ketone and optionally, a leuco dye, a polymerizable monomer or inert components such as binders, solvents and the like are photo-activated in the visible light wavelengths.

IT 19226-99-4

(photosensitizer, for photoactivatable compns. containing hexaarylbiimidazole)

RN 19226-99-4 USPATFULL

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ΑN
       86:35649 USPATFULL
       Process for producing equidensity images using photohardenable materials
ΤI
       Grossa, Mario, Dreieich, Germany, Federal Republic of
ΤN
       E. I. Du Pont de Nemours and Company, Wilmington, DE, United States
PΑ
       (U.S. corporation)
       US 4595651
                               19860617
PΙ
       US 1984-647809
                               19840906 (6)
ΑI
PRAI
       DE 1983-3335309
                           19830929
DT
       Utility
FS
       Granted
EXNAM Primary Examiner: Kittle, John E.; Assistant Examiner: Hamilton, Cynthia
CT.MN
       Number of Claims: 6
ECL
       Exemplary Claim: 1,2
DRWN
       2 Drawing Figure(s); 2 Drawing Page(s)
LN.CNT 460
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       The invention is illustrated by the following examples wherein the
       percentages are by weight. The molecular weights of polymeric compounds
       are weight average molecular weights (Mw). The Mw of
       the polymers can be determined by using a light scattering technique
       using known standard samples, e.g., polystyrene, polymethacrylic acid,
       polymethylmethacrylate, etc. as known to those skilled in the art.
DETD
Copolymer with an .sup. -- Mw of ca. 50,000 comprising
40% tert.-octyl acrylamide,
                           53.6%
25% methyl methacrylate,
15% hydroxypropyl methacrylate,
16% acrylic acid and
 4% tert.-butyl aminoethyl methacrylate
Terpolymer with an .sup.--Mw of ca. 260,000 comprising
56% ethyl acrylate
                           3.8%
37% methyl methacrylate and
 7% acrylic acid
Trimethylol propane triacrylate
Triethylene glycol dimethacrylate
Tetraethyleneglycol dimethacrylate
                           4.0%
Dimer of 2-(o-chlorophenyl)-3,4-diphenyl imidazole
                           5.0%
Compound A from Example 1 1.9%
Leuco Crystal Violet, Basic Violet 3,
C.I. No. 42553
Compound B from Example 1 0.35%
Victoria Pure Blue BO, C.I. No. 42595
                           0.1%
(C.I. Basic Blue 7)
DETD
       Terpolymer with a Mw of ca. 260,000 comprising
DETD
56% ethyl acrylate
                           20.9%
37% methyl methacrylate and
 7% acrylic acid
Carbon black
                           17.1%
Styrene/maleic acid copolymer (1:1),
                           33.7%
partially esterified with sec. butanol
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L43 ANSWER 1 OF 2 USPATFULL on STN

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(-- Mw ca. 10,000)
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Triethylene glycol dimethacrylate

Trimethylol propane triacrylate

8.5%

6-nitroveratraldehyde

1.8%

Dimer of 2-(o-chlorophenyl)-3,4-diphenyl imidazole

8.6%

Compound C

0.9%

Compound C

##STR3##

IT 77-99-6D, ethoxylated, triacrylate 79-10-7D, ester with ethoxylated trimethylolpropane 109-16-0 109-17-1 603-48-5 1707-68-2

2382-96-9 3290-92-4 9003-20-7 9004-95-9 9011-14-7 19226-99-4 20357-25-9 21829-25-4 25135-39-1 56646-84-5

80867-06-7 67016-70-0 96024-63-4

(photoimaging compns. containing, for contour or equidensity images)

IT19226-99-4

(photoimaging compns. containing, for contour or equidensity images)

RN 19226-99-4 USPATFULL

CN Cyclopentanone, 2,5-bis[[4-(dimethylamino)phenyl]methylene]- (9CI) INDEX NAME)

L43 ANSWER 2 OF 2 USPATFULL on STN

72:16254 USPATFULL ΔN

TIHEXAARYLBIIMIDAZOLE BIS (p-DIALKYL-AMINOPHENYL- α , β -UNSATURATED) KETONE COMPOSITIONS

TN Baum, Martin D., Wilmington, DE, United States Henry, Jr., Cyrus P., Wilmington, DE, United States

PA E. I du Pont de Nemours and Company, Wilmington, DE, United States

PT US 3652275 19720328 19700709 (5)

US 1970-53686 DTUtility

FS Granted

AΙ

EXNAM Primary Examiner: Torchin, Norman G.; Assistant Examiner: Fichter, Richard E.

LREP Powell; John R.

CLMN Number of Claims: 36

No Drawings DRWN

LN.CNT 1249

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

a. Films prepared from formulations C, D, F and G were exposed to light DETD of intensity 1.5 mw/cm..sup.2 from a mercury-vapor lamp; two Corning filters 7-54 and one 0-52 filter were used to give 40 mm bands of incident light centered near 366 μ . Under these conditions, formulation F was fully photopolymerized in 8 seconds, formulation G in 16 seconds, illustrating that hexaarylbiimidazole of F is approximately 2 times faster than that of G.

DETD b. Repeating (a), but with light of intensity 10.0 mw /cm..sup.2 (mercury-vapor lamp) with a wavelength range of about 40 mμ centered near 430 mμ, resulting from the use of one Corning 7-59 and one 3-74 filter, gave entirely different results. Under these conditions, formulation D exhibited a photopolymerization rate (16 sec.) about four times that of G (64 sec.). Similarly, the ketone sensitized formulation C is about three times faster than the unsensitized formulation F. These data illustrate the efficacy of ketone IV in photopolymerization.

DETD

c. Repeating (b), including film formulation B, but with light of intensity 25.0 mw/cm..sup.2 and at wavelength greater than 430 $m\mu$, resulting from the use of one Corning 3-72 filter and one I-69 filter, gave still different results. Under these conditions, formulations F and G (no sensitizer) showed no photopolymerization even after 4-minute exposures. Formulations C and D, on the other hand, exhibited photopolymerization rates as shown in 2a, namely 8 and 16 seconds, respectively. Under these long wavelength irradiation conditions, there is, apparently, no absorption of the biimidazole, hence no photodissociation or photo-induced polymerization. The presence of ketone IV, on the other hand, provides a photopolymerization rate equal to irradiation with near ultraviolet light (a).

DETD

"Mylar" (1mil thick) polyester film was coated to a wet thickness of 6 mil using an acetone solution of cellulose acetate butyrate (13.2 grams), triethyleneglycol dimethacrylate (12.5 milliliters), 2,2'-bis(o-chlorophenyl)-4,4',5,5'-tetrakis-(m-methoxyphenyl)biimidazole (3.0 grams), 2-mercaptobenzoxazole (0.10 grams) and various amounts of Michler's ketone (MK), p,p-bis(dimethylamino)benzophenone, and/or ketone sensitizers of this invention, and laminated as in Example 2. The films were irradiated at two different wavelengths, obtained by the use of suitable filters. Irradiation with light at about 366 m μ , incident intensity of 1.00 mw/cm..sup.2, was obtained using two Corning 7-54 filters and one 0-52 filter. Irradiation at 430 m μ , incident intensity of 10.0 mw/cm..sup.2, was obtained using one Corning 7-59 and one 3-74 filter. The irradiation time required to give complete photopolymerization at the two wavelengths is shown in Table III. ##SPC4##

DETD

Incident Intensity

Film 5 mw./cm..sup.2

15 mw./cm..sup.2

26 mw./cm..sup.2

Unsensitized Control		0.06	0.06	
Control with				
Sensitizer II	0.23	0.39	0.53	
Control with				
Sensitizer VIII	0.25	0.41	0.49	

DETD the formulations were coated on 3 mil polyester film, dried; and then laminated with a 1.42 mil polyester film. Exposure of films A, B, and C to light of 1.0 mw/cm..sup.2 intensity from a mercury-vapor lamp resulted in all three films being completely polymerized with the same one second exposure. The light was of broad wavelength centered near 366 m μ via use of two Corning 7-54 and one 0-52 filters.

DETD However, exposure of films A, B and C to light of 10.0 mw /cm..sup.2 intensity but centered near 430 mµ wavelength via use of one Corning 7-59 and one 3-74 filter showed that both films B and C with sensitizers IV and VII gave complete polymerization with 1/4 the exposure time required for the unsensitized film, A. Under these conditions sensitizer VII was as effective as sensitizer IV and it had less background color.

DETD Solution B comprised 0.15 g. of 1,1-dimethyl-3,5-diketocyclohexanene in 10 ml. of methanol. Formulations 1-4 were applied to 3 mil Mylar polyester film at a wet thickness of 5 mil and dried with a heat lamp. The dried films were laminated with a cover sheet of 1 mil Mylar polyester film. The films at a temperature of 75° C., were exposed to a XB0150W1 xenon arc lamp through a Corning 3-72 filter. This

filter passes light of wavelengths greater than 430 m μ . The irradiance at the film plane was 12.5 mw/cm.sup.2. After this exposure the films were exposed to a single flash from a xenon flashtube (Model K, Hico Corporation, Watertown, Mass.) and the resulting optical densities measured with a Macbeth Quantalog transmission densitometer.

IT **19226-99-4** 38394-50-2 38394-52-4 38394-53-5

(photosensitizer, for photoactivatable compns. containing hexaarylbiimidazole)

IT 19226-99-4

(photosensitizer, for photoactivatable compns. containing hexaarylbiimidazole)

RN 19226-99-4 USPATFULL